C AND C BY CLEAN AND CLEAR BLACK OUT BLACKHEAD CLEARING COFFEE SCRUB- salicylic acid lotion

Johnson & Johnson Consumer Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

C & C by Clean & Clear black out blackhead clearing coffee scrub

Drug Facts

Active ingredient

Salicylic Acid (2%)

Purpose

Acne medication

Use

- for the treatment of acne
- clears blackheads

Warnings

For external use only

When using this product

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time
- avoid contact with eyes. If contact occurs, flush thoroughly with water

Keep out of reach of children. if swallowed, get medical help or contact a Poison Control Center right away

Directions

massage onto damp face, avoiding eyes. rinse thoroughly. use daily.

Other Information

store at room temperature

Inactive ingredients

Water, Cocamidopropyl Hydroxysultaine, Sodium C14-16 Olefin Sulfonate, Sorbitol, Cellulose, Sodium Hydrolyzed Potato Starch Dodecenylsuccinate, Acrylates Crosspolymer-4, Fragrance, Sodium Hydroxide, Microcrystalline Wax, Menthol, Disodium EDTA, C12-15 Alkyl Lactate, Benzalkonium Chloride, Cocamidopropyl PG-Dimonium Chloride Phosphate

Questions?

call toll-free 1-877-754-6411 or 215-273-8755 (collect)

Distributed by:

Johnson & Johnson Consumer Inc. Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 124 mL Tube Label

C&C by Clean & Clear

black out

blackhead clearing coffee scrub salicylic acid acne medication

4.2 fl oz (124mL)



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salicylic acid lotion

Product Information	uct Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0346		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Salicylic Acid (UNII: O414PZ4LPZ) (Salicylic Acid - UNII:O414PZ4LPZ)	Salicylic Acid	20 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
Water (UNII: 059QF0KO0R)		
Cocamidopropyl Hydroxysultaine (UNII: 62V75NI93W)		
Sodium C14-16 Olefin Sulfonate (UNII: O9W3D3YF5U)		
Sorbitol (UNII: 506T60A25R)		
Powdered Cellulose (UNII: SMD1X3XO9M)		
Edetate Disodium Anhydrous (UNII: 8NLQ36F6MM)		
C12-15 Alkyl Lactate (UNII: GC844VRD7E)		
Benzalkonium Chloride (UNII: F5UM2KM3W7)		
Sodium Hydroxide (UNII: 55X04QC32I)		
Cocamidopropyl Propylene Glycol-Dimonium Chloride Phosphate (UNII: H2KVQ74JM4)		
Microcrystalline Wax (UNII: XOF597Q3KY)		
Menthol, Unspecified Form (UNII: L7T10 EIP3A)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69968-0346-4	124 mL in 1 TUBE; Type 0: Not a Combination Product	06/01/2018	05/31/2021	
2	NDC:69968-0346-1	30 mL in 1 TUBE; Type 0: Not a Combination Product	06/01/2018	05/31/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part333D	06/01/2018	05/31/2021	

Labeler - Johnson & Johnson Consumer Inc. (002347102)

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